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# B. Braun Vet Care hypertonic NaCl-Solution (75 mg/ml) Solution for infusion for horses, cattle, sheep, goats, pigs, dogs and cats

Authorised

- Sodium chloride

## Product identification

**Medicine name:**
B. Braun Vet Care hypertonic NaCl-Solution (75 mg/ml) Solution for infusion for horses, cattle, sheep, goats, pigs, dogs and cats

**Active substance:**
Sodium chloride

**Target species:**
Cattle
Sheep
Goat
Horse
Pig
Dog
Cat

**Route of administration:**
Intravenous use

## Product details

**Active substance and strength:**
Sodium chloride
7.50 gram(s) / 100.00 millilitre(s)

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**Pharmaceutical form:**
Solution for infusion

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**Withdrawal period by route of administration:**
**Intravenous use:**

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Horse**
  - Meat and offal. 0 day
  - Milk. 0 day
- **Pig**
  - Meat and offal. 0 day
  - Milk. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QB05BB

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
France

**Available in:**
France

**Package description:**
box containing 1 bottle of 500 ml
box containing 10 bottles of 500 ml

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Well-established use application (Article 13a of Directive No 2001/82/EC)

**Marketing authorisation holder:**
B. Braun Melsungen AG

**Marketing authorisation date:**
10/09/2009

**Manufacturing sites for batch release:**
B BRAUN MEDICAL S.A.

**Responsible authority:**
French Agency For Food, Environmental And Occupational Health & Safety

**Authorisation number:**

FR/V/9398027 6/2009

**Date of authorisation status change:**
10/09/2014

**Reference member state:**
Spain

**Procedure number:**
ES/V/0149/001

**Concerned member states:**
Austria Belgium Denmark France Germany Italy Netherlands Portugal Romania Sweden United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Summary of Product Characteristics

English (PDF)
Published on: 25/06/2025
Updated on: 13/03/2026
Download

Labelling

Package Leaflet

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

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